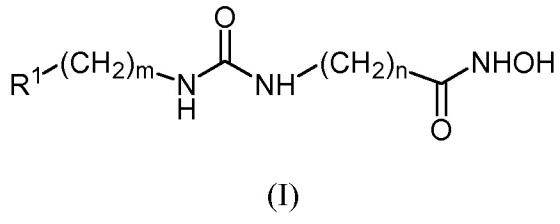


Claims:

1. **(Previously presented)** A compound having the formula



or a pharmaceutically acceptable salt thereof,

wherein

R^1 is aryl, -C₃-C₇ cycloalkyl, adamantyl, or -3- to 10-membered heterocycle, any of which may be unsubstituted or substituted with one or more of the following groups: -halo, -C₁-C₆ alkyl, -O-(C₁-C₆ alkyl), -OH, -CN, -COOR', -OC(O)R', NHR', N(R')₂, -NHC(O)R' or -C(O)NHR' groups wherein R' is -H or unsubstituted -C₁-C₆ alkyl, with the proviso that when n is 2, R^1 cannot be -C₃-C₇ cycloalkyl or -3- to 10-membered heterocycle,

m is an integer ranging from 1-10; and

n is an integer ranging from 1-10.

2. **(Original)** The compound of claim 1 wherein R^1 is phenyl.

3. **(Original)** The compound of claim 1 wherein n is an integer ranging from 1-5.

4. **(Original)** The compound of claim 1 wherein m is 2.

5. **(Original)** The compound of claim 1 wherein R^1 is phenyl, m is 2 and n is 3.

6. **(Original)** The compound of claim 1 wherein R^1 is -4-N(CH₃)₂-phenyl and m is 1.

7. **(Original)** The compound of claim 1 wherein R^1 is -4-N(CH₃)₂-phenyl, m is 1 and n is 4.

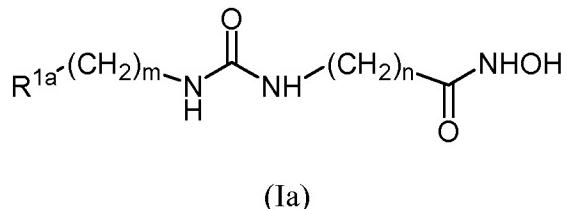
8. **(Original)** The compound of claim 1 wherein R^1 is -4-N(CH₃)₂-phenyl, m is 1 and n is 5.

Claims 9 - 31 (**canceled**)

32. **(Original)** A pharmaceutical composition comprising the compound or a pharmaceutically acceptable salt of the compound of claim 1 and a pharmaceutically acceptable carrier or vehicle.

Claims 33 - 40 (**canceled**)

41. (**Withdrawn**) A method for increasing the sensitivity of a cancer cell to the cytotoxic effects of radiotherapy, said method comprising contacting said cell with a compound having the formula:



or a pharmaceutically acceptable salt thereof,

wherein

R^{1a} is aryl, $-\text{C}_3\text{-C}_7$ cycloalkyl, adamantyl, or $-3\text{- to } 10$ -membered heterocycle, any of which may be unsubstituted or substituted with one or more of the following groups: -halo, $-\text{C}_1\text{-C}_6$ alkyl, $-\text{O}-(\text{C}_1\text{-C}_6 \text{ alkyl})$, $-\text{OH}$, $-\text{CN}$, $-\text{COOR}'$, $-\text{OC(O)R}'$, NHR' , $\text{N}(\text{R}')_2$, $-\text{NHC(O)R}'$ or $-\text{C(O)NHR}'$ groups wherein R' is $-\text{H}$ or unsubstituted $-\text{C}_1\text{-C}_6$ alkyl;

m is an integer ranging from 0-10; and

n is an integer ranging from 1-10,

in an amount sufficient to increase the sensitivity of said cell to the cytotoxic effects of radiotherapy.

Claims 42-49 (**Canceled**).

50. (**Withdrawn**) The method of claim 41 wherein the cell is an *in vivo* cell.

51. (**Withdrawn**) A method for treating cancer, said method comprising administering to a subject in need thereof the compound or a pharmaceutically acceptable salt of the compound of claim 1 in an amount sufficient to treat said cancer.

Claims 52-59 (**Canceled**).

60. (**Withdrawn**) The method of claim 51 wherein the subject is a human.

61. (**Withdrawn**) The method of claim 51 wherein the cancer is Non-Hodgkin's lymphoma, Hodgkin's disease, Ewing's sarcoma, testicular cancer, prostate cancer, larynx cancer, cervical

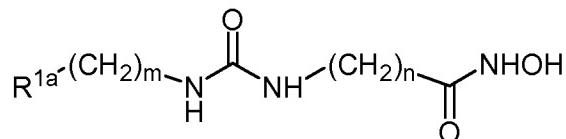
cancer, nasopharynx cancer, breast cancer, colon cancer, pancreatic cancer, head and neck cancer, esophageal cancer, rectal cancer, small-cell lung cancer, non-small cell lung cancer, brain cancer, or a CNS neoplasm.

62. **(Withdrawn)** The method of claim 51 further comprising administering to said subject another therapeutic agent or a pharmaceutically acceptable salt thereof.

63. **(Withdrawn)** The method of claim 62 wherein the other therapeutic agent is an anticancer agent.

64. **(Withdrawn)** A method for treating cancer, said method comprising:

(a) administering to a subject in need thereof, a compound having the formula:



(Ia)

or a pharmaceutically acceptable salt thereof,

wherein

R^{1a} is aryl, -C₃-C₇ cycloalkyl, adamantly or -3- to 10-membered heterocycle, any of which may be unsubstituted or substituted with one or more of the following groups: -halo, -C₁-C₆ alkyl, -O-(C₁-C₆ alkyl), -OH, -CN, -COOR', -OC(O)R', NHR', N(R')₂, -NHC(O)R' or -C(O)NHR' groups wherein R' is -H or unsubstituted -C₁-C₆ alkyl;

m is an integer ranging from 0-10; and

n is an integer ranging from 1-10,

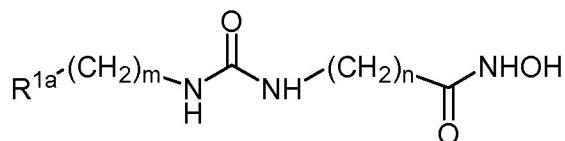
in an amount sufficient to sensitize a cancer cell to the cytotoxic effects of radiotherapy; and

(b) administering to said subject an amount of radiotherapy sufficient to treat said cancer.

Claims 65 - 72 **(Cancelled)**.

73. **(Withdrawn)** The method of claim 64 wherein the compound administered in step (a) and the radiotherapy administered in step (b) act adjunctively.

74. **(Withdrawn)** The method of claim 64 wherein the subject is a human.
75. **(Withdrawn)** The method of claim 64 wherein the cancer is Non-Hodgkin's lymphoma, Hodgkin's disease, Ewing's sarcoma, testicular cancer, prostate cancer, larynx cancer, cervical cancer, nasopharynx cancer, breast cancer, colon cancer, pancreatic cancer, head and neck cancer, esophageal cancer, rectal cancer, small-cell lung cancer, non-small cell lung cancer, brain cancer, or a CNS neoplasm.
76. **(Withdrawn)** The method of claim 64 further comprising administering to said subject another therapeutic agent or a pharmaceutically acceptable salt thereof.
77. **(Withdrawn)** The method of claim 76 wherein the other therapeutic agent is an anticancer agent.
78. **(Withdrawn)** The method of claim 64 wherein the administering of step (a) is done prior to the administering of step (b).
79. **(Withdrawn)** The method of claim 64 wherein the administering of step (a) is done subsequent to the administering of step (b).
80. **(Withdrawn)** The method of claim 64 wherein the administering of step (a) and the administering of step (b) are done concurrently.
81. **(Withdrawn)** A method for treating a neurological disease, said method comprising administering to a subject in need thereof a compound having the formula



(Ia)

or a pharmaceutically acceptable salt thereof,

wherein

$\text{R}^{1a'}$ is aryl, -C₃-C₇ cycloalkyl, adamantyl, or -3- to 10-membered heterocycle, any of which may be unsubstituted or substituted with one or more of the following groups: -halo, -C₁-C₆ alkyl, -

$O-(C_1-C_6 \text{ alkyl})$, $-OH$, $-CN$, $-COOR'$, $-OC(O)R'$, NHR' , $N(R')_2$, $-NHC(O)R'$ or $-C(O)NHR'$ groups wherein R' is $-H$ or unsubstituted $-C_1-C_6 \text{ alkyl}$;

m is an integer ranging from 0-10; and

n is an integer ranging from 1-10,

in an amount sufficient to treat said neurological disease.

Claims 82 - 89 (**Canceled**).

90. (**Withdrawn**) The method of claim 81 wherein said disease of the central nervous system is Huntington's disease, lupus, or schizophrenia.

91. (**Withdrawn**) The method of claim 81 wherein the subject is a human.

92. (**Previously presented**) The compound of claim 1 wherein R^1 is $-4-N(CH_3)_2\text{-phenyl}$, m is 1 and n is 6.

93. (**Previously presented**) The compound of claim 1 wherein R^1 is $-4-N(CH_3)_2\text{-phenyl}$, m is 1 and n is 7.

94. (**New**) The compound of claim 1, wherein R^1 is phenyl or adamantyl, either of which may be unsubstituted or substituted with one or more of the following groups: $-halo$, $-C_1-C_6 \text{ alkyl}$, $-O-(C_1-C_6 \text{ alkyl})$, $-OH$, $-CN$, $-COOR'$, $-OC(O)R'$, NHR' , $N(R')_2$, $-NHC(O)R'$ or $-C(O)NHR'$ groups wherein R' is $-H$ or unsubstituted $-C_1-C_6 \text{ alkyl}$.